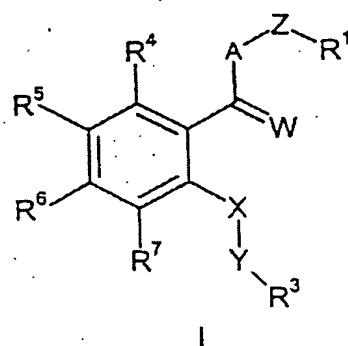
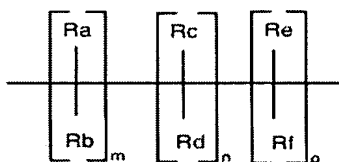


1. A compound of formula I

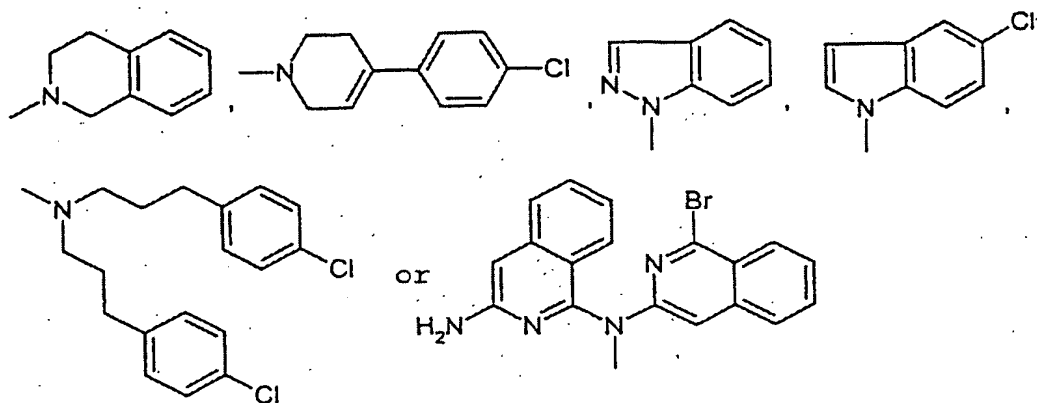


in which

- A stands for the group =NR²,
W stands for oxygen, sulfur, two hydrogen atoms or the group =NR⁸,
Z stands for the group =NR¹⁰ or =N-,
-N(R¹⁰)-(CH₂)_q-, branched or unbranched C₁₋₆ alkyl or the group



or A, Z and R¹ together form the group



A2

m, n and o

stand for 0-3,

q

stands for 1-6,

R_a, R_b, R_c, R_d, R_e, R_f,

independently of one another, stand for hydrogen, C₁₋₄ alkyl or the group =NR¹⁰, and/or R_a and/or R_b can form a bond with R_c and/or R_d or R_c can form a bond with R_e and/or R_f or up to two of radicals R_a-R_f can close a bridge with up to 3 C-atoms each to form R¹ or R²,

X

stands for the group =NR⁹ or =N-,

Y

stands for the group -(CH₂)_p,

p

stands for 1-4,

R¹

stands for unsubstituted aryl or heteroaryl, or for aryl or heteroaryl substituted one or more times with halogen; C₁₋₆ alkyl; or one or more times with halogen substituted C₁₋₆ alkyl or C₁₋₆ alkoxy; with the proviso that R¹ is not aryl directly bonded to =NR² in the meaning of A,

R²

stands for hydrogen or C₁₋₆ alkyl or forms a bridge with up to 3 ring members with R_a-R_f from Z or to form R₁,

R³

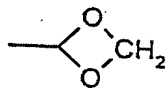
stands for monocyclic or bicyclic aryl or heteroaryl that is unsubstituted or optionally substituted in one or more places with halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy or hydroxy,

R⁴, R⁵, R⁶, and R⁷,

independently of one another, stand for hydrogen, halogen, or C₁₋₆ alkoxy, C₁₋₆ alkyl or C₁₋₆ carboxylalkyl that is unsubstituted or optionally substituted in one or more places with halogen,

or R⁵ and R⁶ together form the group

A²



R⁸, R⁹, and R¹⁰,

independently of one another, stand for hydrogen or C₁₋₆ alkyl,

or an isomer or pharmaceutically acceptable salt thereof,

with the proviso that when A is =NR², X is =NR⁹, R^{2,4,6,7,9} is H, R⁵ is Cl, W is O, Z=Y is -CH₂-, and R³ is 4-pyridyl, then R¹ is not 3,4-methylenedioxybenzyl. *Directed to Example 3*

7. (Amended) A method of claim 11 for the treatment of tumors, psoriasis, arthritis, hemangioma, angiofibroma, eye diseases, neovascular glaucoma, renal diseases, fibrotic diseases, mesangial-cell-proliferative diseases, arteriosclerosis, injuries to the nerve tissue, and for inhibiting the reocclusion of vessels after balloon catheter treatment, in vascular prosthetics or after mechanical devices are used to keep vessels open.

A³

8. (Amended) A pharmaceutical composition comprising a therapeutical effective amount of at least one compound according to claim 1 and a pharmaceutical acceptable carrier.

9. (Amended) A pharmaceutical composition according to claim 8 for the treatment of tumors, psoriasis, arthritis, such as rheumatoid arthritis, hemangioma, angiofibroma, eye diseases, such as diabetic retinopathy, neovascular glaucoma, renal diseases, such as glomerulonephritis, diabetic nephropathy, malignant nephrosclerosis, thrombotic microangiopathic syndrome, transplant rejections and glomerulopathy, fibrotic diseases, such as cirrhosis of the liver, mesangial-cell-proliferative diseases, arteriosclerosis, injuries to the nerve tissue, and for inhibiting the reocclusion of vessels after balloon catheter treatment, in vascular prosthetics or after mechanical devices are used to keep vessels open, such as, e.g., stents..

A⁴

11. (Amended) A method of inhibiting the tyrosine kinase KDR and/or FLT, comprising *enabling* administering to a patient in need thereof a therapeutically effective amount of a compound according to claim 1.

A4

12. (Amended) A method of producing a pharmaceutical preparation for enteral, parenteral and oral administration comprising mixing a compound of claim 1 with a suitable pharmaceutical carrier.
